FDA 510(k) Premarket Notification

K993092 SONADA ULTRASOUND SYSTEM

CLEARED

Apr 14, 2000
211 days to decision

K993092 · Product code: **OXO** · Radiology
Source: https://www.510kdatabase.net/k993092/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Image-intensified Fluoroscopic X-ray System, Mobile (OXO)
Date received	Sep 16, 1999
Decision date	Apr 14, 2000
Days to decision	211 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Sonovision Corp.**
Location	Hasbrouck Heights, NJ, US
Contact	PAUL SCHNEIDER
510(k) history	1 submissions · 1 cleared · 2000-2000